Claim Amendments

1. **(previously presented)** A pharmaceutical composition for intramammary administration to a non-human mammal, wherein:

the composition comprises:

an antibacterial agent,

prednisolone, and

a pharmaceutically acceptable carrier; and

the composition comprises at least 20 mg of prednisolone per unit dose.

2. **(previously presented)** The composition according to claim 1, wherein the composition comprises prednisolone in an amount of 20 to 40 mg per unit dose.

3. **(previously presented)** The composition according to claim 2, wherein the composition comprises prednisolone in an amount of 20 to 30 mg per unit dose.

4. **(previously presented)** The composition according to claim 1, wherein the antibacterial agent is a cephalosporin.

5. **(previously presented)** The composition according to claim 4, wherein the cephalosporin is cephapirin.

6. **(previously presented)** The composition according to claim 4, wherein the cephalosporin is cefquinome.

7. **(previously presented)** The composition according to claim 1, wherein the composition comprises the antibacterial agent in an amount of 10 to 500 mg per unit dose.

8. **(withdrawn)** A process for preparing a pharmaceutical composition according to claim 1, comprising the steps of mixing an oil and one or more pharmaceutically acceptable

additives to form a carrier, and suspending the antibacterial agent and the prednisolone in the carrier.

9. (Canceled).